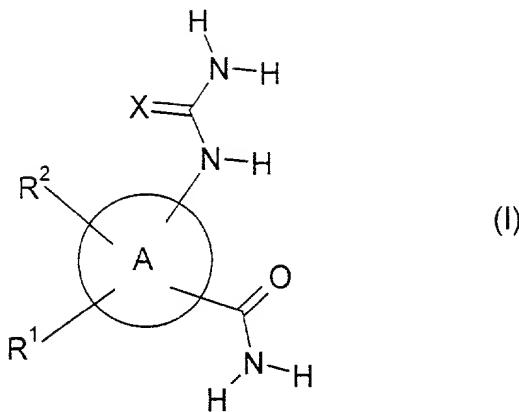


CLAIMS

1. A compound of formula (I)



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in which:

A represents a 5-membered heteroaromatic ring containing one or two heteroatoms

10 selected independently from oxygen, nitrogen or sulfur;

R¹ represents a phenyl group or a 5- to 7-membered heteroaromatic ring containing one to three heteroatoms selected independently from oxygen, nitrogen or sulfur; said phenyl or heteroaromatic ring being optionally substituted by one or more substituents selected
 15 independently from halogen, cyano, nitro, -NR³R⁴, -CONR⁵R⁶, -COOR⁷, -NR⁸COR⁹, -SR¹⁰, -S(O)_mR¹⁰, -SO₂NR⁵R⁶, -NR⁸SO₂R¹⁰, C₁-C₆ alkyl, trifluoromethyl, -(CH₂)_nR¹¹, -O(CH₂)_nR¹¹ or -OR¹²;

R² represents hydrogen, halogen, cyano, nitro, -NR¹³R¹⁴, -CONR¹⁵R¹⁶, -COOR¹⁷,
 20 -NR¹⁸COR¹⁹, -S(O)_mR²⁰, -SO₂NR¹⁵R¹⁶, -NR¹⁸SO₂R²⁰, C₁-C₂ alkyl, trifluoromethyl, C₂-C₃ alkenyl, C₂-C₃ alkynyl, trifluoromethoxy, C₁-C₂ alkoxy or C₁-C₂ alkanoyl;

X represents oxygen or sulphur;

each of R³, R⁴, R⁵, R⁶, R⁷, R⁸, R⁹, R¹⁰ and R¹² independently represent a hydrogen atom or C₁-C₆ alkyl;

R¹¹ represents NR²¹R²² where R²¹ and R²² are independently hydrogen or C₁-C₆ alkyl

5 optionally substituted by C₁-C₄ alkoxy; or R²¹ and R²² together with the nitrogen atom to which they are attached form a 5- or 6-membered saturated ring optionally containing a further O, S or NR²³ group where R²³ is hydrogen or C₁-C₆ alkyl; or R¹¹ represents OR²⁴ where R²⁴ represents C₁-C₆ alkyl;

10 each of R¹³, R¹⁴, R¹⁵, R¹⁶, R¹⁷, R¹⁸, R¹⁹ and R²⁰ independently represent a hydrogen atom or C₁-C₂ alkyl;

m represents an integer 0, 1 or 2;

15 n represents an integer 2, 3 or 4;

and optical isomers, racemates and tautomers thereof and pharmaceutically acceptable salts or solvates thereof:

20 provided that:

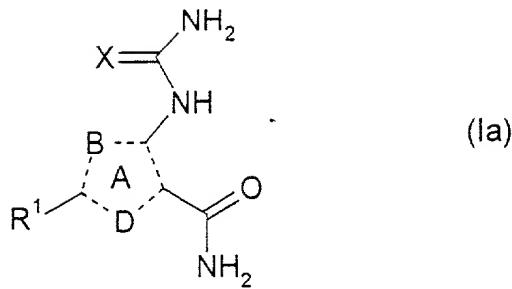
when A represents thiophene, furan or pyrrole, then R¹ is not 4-pyridinyl or 3-pyrazolyl; and

when A represents oxazole, thiazole or imidazole, then R¹ is not 3-pyridinyl or 5-pyrimidyl.

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2. A compound of formula (I), according to Claim 1, wherein X represents oxygen.

3. A compound of formula (I), according to Claim 1 or Claim 2, in which the group A is substituted as shown below in formula (Ia), where B and D are selected from CR², S, O and NR²⁵, where R² is as defined in Claim 1 and R²⁵ is hydrogen or C₁-C₆ alkyl:



4. A compound according to any one of Claims 1 to 3 in which the ring A is thiophene.

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5. A compound according to any one of Claims 1 to 4 in which R¹ represents optionally substituted phenyl.

6. A compound according to any one of Claims 1 to 5 in which R² represents H or

10 methyl.

7. A compound according to Claim 6 in which R² represents H.

8. A compound of formula (I), according to claim 1, selected from:

15 3-[(aminocarbonyl)amino]-5-phenyl-2-thiophenecarboxamide;

3-[(aminocarbonyl)amino]-5-(3-chlorophenyl)-2-thiophenecarboxamide;

3-[(aminocarbonyl)amino]-5-(4-fluorophenyl)-2-thiophenecarboxamide;

3-[(aminocarbonyl)amino]-5-(4-chlorophenyl)-2-thiophenecarboxamide;

3-[(aminocarbonyl)amino]-5-(4-isobutylphenyl)-2-thiophenecarboxamide;

20 3-[(aminocarbonyl)amino]-5-(2-thienyl)-2-thiophenecarboxamide;

3-[(aminocarbonyl)amino]-5-(4-methoxyphenyl)-2-thiophenecarboxamide;

3-[(aminocarbonyl)amino]-5-(3-thienyl)-2-thiophenecarboxamide;

3-[(aminocarbonyl)amino]-5-(3-hydroxyphenyl)-2-thiophenecarboxamide;

3-[(aminocarbonyl)amino]-5-(2-chlorophenyl)-2-thiophenecarboxamide;

3-[(aminocarbonyl)amino]-5-(2-methoxyphenyl)-2-thiophenecarboxamide;

3-[(aminocarbonyl)amino]-5-{2-[2-(dimethylamino)ethoxy]phenyl}-2-thiophenecarboxamide;

5 3-[(aminocarbonyl)amino]-5-{4-[2-(dimethylamino)ethoxy]phenyl}-2-thiophenecarboxamide;

3-[(aminocarbonyl)amino]-5-(3-methoxyphenyl)-2-thiophenecarboxamide;

2-[(aminocarbonyl)amino]-5-phenyl-3-thiophenecarboxamide;

3-[(aminocarbonyl)amino]-5-{4-[2-(1-morpholinyl)ethoxy]phenyl}-2-thiophenecarboxamide;

10 thiophenecarboxamide;

3-[(aminocarbonyl)amino]-5-{4-[2-(1-pyrrolidinyl)ethoxy]phenyl}-2-thiophenecarboxamide;

3-[(aminocarbonyl)amino]-5-{4-[2-(1-piperidinyl)ethoxy]phenyl}-2-thiophenecarboxamide;

15 3-[(aminocarbonyl)amino]-5-{4-[3-(dimethylamino)propoxy]phenyl}-2-thiophenecarboxamide;

3-[(aminocarbonyl)amino]-5-{3-[2-(dimethylamino)ethoxy]phenyl}-2-thiophenecarboxamide;

3-[(aminocarbonyl)amino]-5-{3-[2-(1-morpholinyl)ethoxy]phenyl}-2-thiophenecarboxamide;

20 thiophenecarboxamide;

3-[(aminocarbonyl)amino]-5-{3-[2-(1-pyrrolidinyl)ethoxy]phenyl}-2-thiophenecarboxamide;

3-[(aminocarbonyl)amino]-5-{3-[2-(1-piperidinyl)ethoxy]phenyl}-2-

thiophenecarboxamide;

3-[(aminocarbonyl)amino]-5-{3-[3-(dimethylamino)propoxy]phenyl}-2-

thiophenecarboxamide;

5 3-[(aminocarbonyl)amino]-5-{2-[2-(1-morpholinyl)ethoxy]phenyl}-2-

thiophenecarboxamide;

3-[(aminocarbonyl)amino]-5-{2-[2-(1-pyrrolidinyl)ethoxy]phenyl}-2-

thiophenecarboxamide;

3-[(aminocarbonyl)amino]-5-{2-[2-(1-piperidinyl)ethoxy]phenyl}-2-

10 thiophenecarboxamide;

3-[(aminocarbonyl)amino]-5-{2-[3-(dimethylamino)propoxy]phenyl}-2-

thiophenecarboxamide;

2-[(aminocarbonyl)amino]-4-methyl-5-(4-chlorophenyl)-3-thiophenecarboxamide;

2-[(aminocarbonyl)amino]-4-methyl-5-(4-methylphenyl)-3-thiophenecarboxamide;

15 2-[(aminocarbonyl)amino]-4-ethyl-5-phenyl-3-thiophenecarboxamide;

2-[(aminocarbonyl)amino]-4-methyl-5-(4-methoxyphenyl)-3-thiophenecarboxamide;

2-[(aminocarbonyl)amino]-4-methyl-5-(4-fluorophenyl)-3-thiophenecarboxamide;

2-[(aminocarbonyl)amino]-4-methyl-5-(3-fluorophenyl)-3-thiophenecarboxamide;

2-[(aminocarbonyl)amino]-4-methyl-5-(3-methoxyphenyl)-3-thiophenecarboxamide;

20 2-[(aminocarbonyl)amino]-4-methyl-5-(3-chloro-4-methoxyphenyl)-3-

thiophenecarboxamide;

2-[(aminocarbonyl)amino]-4-methyl-5-(2-chlorophenyl)-3-thiophenecarboxamide;

2-[(aminocarbonyl)amino]-4-methyl-5-(3-trifluoromethylphenyl)-3-thiophenecarboxamide;

2-[(aminocarbonyl)amino]-4-methyl-5-(3-methyl-4-methoxyphenyl)-3-thiophenecarboxamide;

2-[(aminocarbonyl)amino]-4-methyl-5-(3,5-dimethoxyphenyl)-3-thiophenecarboxamide;

2-[(aminocarbonyl)amino]-4-methyl-5-(2,3-dimethoxyphenyl)-3-thiophenecarboxamide;

5 2-[(aminocarbonyl)amino]-4-methyl-5-(4-isopropylphenyl)-3-thiophenecarboxamide;

2-[(aminocarbonyl)amino]-4-methyl-5-(3,4,5-trimethoxyphenyl)-3-thiophenecarboxamide;

2-[(aminocarbonyl)amino]-4-methyl-5-(2-pyridyl)-3-thiophenecarboxamide;

2-[(aminocarbonyl)amino]-5-[2-(5-methoxypyridyl)]-4-methyl-3-thiophenecarboxamide;

2-[(aminocarbonyl)amino]-4-methyl-5-(4-pyrimidyl)-3-thiophenecarboxamide;

10 2-[(aminocarbonyl)amino]-4-methyl-5-(2-pyrazinyl)-3-thiophenecarboxamide;

2-[(aminocarbonyl)amino]-4-methyl-5-(3,4-dichlorophenyl)-3-thiophenecarboxamide;

2-[(aminocarbonyl)amino]-4-methyl-5-(4-cyanophenyl)-3-thiophenecarboxamide;

2-[(aminocarbonyl)amino]-4-methyl-5-(4-hydroxyphenyl)-3-thiophenecarboxamide;

2-[(aminocarbonyl)amino]-4-methyl-5-(4-[2-(1-piperidinyl)ethoxy]phenyl)-3-

15 thiophenecarboxamide;

2-[(aminocarbonyl)amino]-4-methyl-5-(4-[2-(diethylamino)ethoxy]phenyl)-3-

thiophenecarboxamide;

2-[(aminocarbonyl)amino]-4-methyl-5-(2-furyl)-3-thiophenecarboxamide;

2-[(aminocarbonyl)amino]-4-trifluoromethyl-5-phenyl-3-thiophenecarboxamide;

20 2-[(aminocarbonyl)amino]-4-methyl-5-(2-(4-methylthiazolyl))-3-thiophenecarboxamide;

2-[(aminocarbonyl)amino]-4-methyl-5-phenyl-3-thiophenecarboxamide;

2-[(aminocarbonyl)amino]-4-methyl-5-(3-methyl-isoxazol-5-yl)-3-thiophenecarboxamide;

2-[(aminocarbonyl)amino]-5-(4-cyanophenyl)-3-thiophenecarboxamide;

2-[(aminocarbonyl)amino]-5-(4-trifluoromethylphenyl)-3-thiophenecarboxamide;

2-[(aminocarbonyl)amino]-5-(2,4-difluorophenyl)-3-thiophenecarboxamide;

2-[(aminocarbonyl)amino]-5-(2-pyridyl)-3-thiophenecarboxamide;

2-[(aminocarbonyl)amino]-5-(3-pyridyl)-3-thiophenecarboxamide;

5 2-[(aminocarbonyl)amino]-5-[5-(2-methoxypyridyl)-3-thiophenecarboxamide;

2-[(aminocarbonyl)amino]-5-[5-(2,4-dimethoxypyrimidyl)-3-thiophenecarboxamide;

2-[(aminocarbonyl)amino]-5-(4-hydroxyphenyl)-3-thiophenecarboxamide;

2-[(aminocarbonyl)amino]-5-(4-chlorophenyl)-3-thiophenecarboxamide;

2-[(aminocarbonyl)amino]-5-(4-methanesulphonylphenyl)-3-thiophenecarboxamide;

10 2-[(aminocarbonyl)amino]-5-(2-(N-t-butoxycarbonyl)pyrrolyl)-3-thiophenecarboxamide;

2-[(aminocarbonyl)amino]-5-(2-(5-cyanothienyl)-3-thiophenecarboxamide;

2-[(aminocarbonyl)amino]-5-(3,5-dimethyl-isoxazol-4-yl)-3-thiophenecarboxamide;

2-[(aminocarbonyl)amino]-5-(3-furyl)-3-thiophenecarboxamide;

2-[(aminocarbonyl)amino]-5-(2-pyrrolyl)-3-thiophenecarboxamide;

15 2-[(aminocarbonyl)amino]-5-(5-pyrimidinyl)-3-thiophenecarboxamide;

2-[(aminocarbonyl)amino]-5-(2-(5-chlorothienyl)-3-thiophenecarboxamide;

2-[(aminocarbonyl)amino]-5-[2-(5-trifluoromethylpyridyl)-3-thiophenecarboxamide;

2-[(aminocarbonyl)amino]-5-[2-(5-bromopyridyl)-3-thiophenecarboxamide;

2-[(aminocarbonyl)amino]-5-(2-(5-cyanofuryl)-3-thiophenecarboxamide;

20 2-[(aminocarbonyl)amino]-5-(4-[2-(1-piperidinyl)ethoxy]phenyl)-3-thiophenecarboxamide;

2-[(aminocarbonyl)amino]-5-(4-[2-(1-(2,2,6,6-tetramethyl)piperidinyl)ethoxy]phenyl)-3-thiophenecarboxamide;

2-[(aminocarbonyl)amino]-5-(4-(thiazol-4-yl-methoxy)phenyl)-3-thiophenecarboxamide;

2-[(aminocarbonyl)amino]-5-(4-[2-(dimethylamino)ethoxy]phenyl)-3-

thiophenecarboxamide;

2-[(aminocarbonyl)amino]-5-(4-[2-(diethylamino)ethoxy]phenyl)-3-

thiophenecarboxamide;

2-[(aminocarbonyl)amino]-5-(4-[2-(1-morpholinyl)ethoxy]phenyl)-3-

thiophenecarboxamide;

2-[(aminocarbonyl)amino]-5-(2-furyl)-3-thiophenecarboxamide;

2-[(aminocarbonyl)amino]-5-(2-(5-methylfuryl))-3-thiophenecarboxamide;

10 5-[(aminocarbonyl)amino]-2-(3,5-dichlorophenyl)-1,3-oxazole-4-carboxamide;

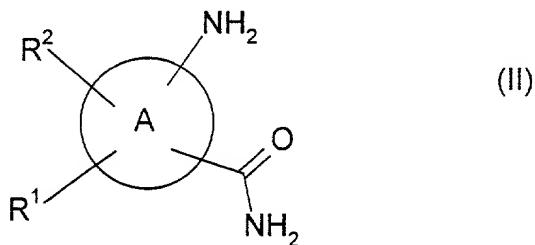
5-[(aminocarbonyl)amino]-2-(4-trifluoromethylphenyl)-1,3-oxazole-4-carboxamide;

2-[(aminothiocarbonyl)amino]-5-phenyl-3-thiophenecarboxamide;

and pharmaceutically acceptable salts and solvates thereof.

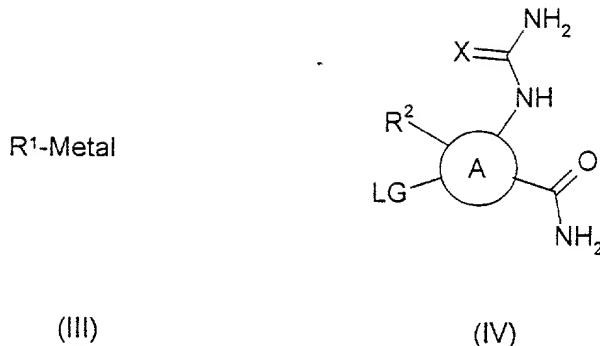
15 9. A process for the preparation of a compound of formula (I), according to any one of Claims 1 to 8, which comprises:

(a) reaction of a compound of formula (II):



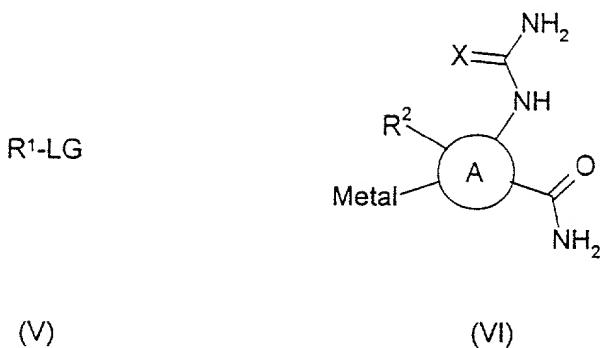
20 wherein A, R¹ and R² are as defined in Claim 1 with an isocyanate (X = O) or an isothiocyanate (X = S); or

(b) reaction of compound of formula (III) with a compound of formula (IV)



wherein A, X, R¹ and R² are as defined in Claim 1 and LG represents a leaving group; or

5 (c) reaction of compound of formula (V) with a compound of formula (VI)



wherein A, X, R¹ and R² are as defined in Claim 1 and LG represents a leaving group;

10 and where necessary converting the resultant compound of formula (I), or another salt thereof, into a pharmaceutically acceptable salt thereof; or converting the resultant compound of formula (I) into a further compound of formula (I); and where desired converting the resultant compound of formula (I) into an optical isomer thereof.

15 10. A pharmaceutical composition comprising a compound of formula (I), or a pharmaceutically acceptable salt or solvate thereof, as claimed in any one of claims 1 to 8 in association with a pharmaceutically acceptable adjuvant, diluent or carrier.

11. A process for the preparation of a pharmaceutical composition as claimed in Claim 20 10 which comprises mixing a compound of formula (I), or a pharmaceutically acceptable

salt or solvate thereof, as claimed in any one of claims 1 to 8 with a pharmaceutically acceptable adjuvant, diluent or carrier.

12. A compound of formula (I), or a pharmaceutically-acceptable salt or solvate thereof, as

5 claimed in any one of claims 1 to 8 for use in therapy.

13. Use of a compound of formula (I), or a pharmaceutically acceptable salt or solvate

thereof, as claimed in any one of claims 1 to 8 in the manufacture of a medicament for use

in therapy.

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14. Use of a compound of formula (I), or a pharmaceutically acceptable salt or solvate

thereof, as claimed in any one of claims 1 to 8 in the manufacture of a medicament for use

in the treatment or prophylaxis of diseases or conditions in which inhibition of IKK2

activity is beneficial.

15

15. Use of a compound of formula (I), or a pharmaceutically acceptable salt or solvate

thereof, as claimed in any one of claims 1 to 8 in the manufacture of a medicament for use

in the treatment or prophylaxis of inflammatory disease.

20

16. The use as claimed in Claim 15 wherein the disease is asthma.

17. The use as claimed in Claim 15 wherein the disease is rheumatoid arthritis.

18. The use as claimed in Claim 15 wherein the disease is multiple sclerosis.

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19. The use as claimed in Claim 15 wherein the disease is chronic obstructive

pulmonary disease.

20. A method of treating an IKK2 mediated disease which comprises administering to a

30 patient a therapeutically effective amount of a compound of formula (I), or a

pharmaceutically acceptable salt or solvate thereof, as claimed in any one of claims 1 to 8.

21. A method of treating an inflammatory disease in a patient suffering from, or at risk of, said disease, which comprises administering to the patient a therapeutically effective amount of a compound of formula (I), or a pharmaceutically acceptable salt or solvate thereof, as claimed in any one of claims 1 to 8.

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22. A method according to claim 21, wherein the disease is asthma.

23. A method according to claim 21, wherein the disease is rheumatoid arthritis.

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24. A method according to claim 21, wherein the disease is multiple sclerosis.

25. A method according to claim 21, wherein the disease is chronic obstructive pulmonary disease.